

Vitofyllin 50 mg film-coated tablets for dogs

Authorised

- Propentofylline

Product identification

Medicine name:

Vitofyllin 50 mg film-coated tablets for dogs
Vitofyllin 50 mg filmom obalené tablety pre psy

Active substance:

Propentofylline

Target species:

Dog

Route of administration:

Oral use

Product details

Active substance and strength:

Propentofylline
50.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:

Film-coated tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QC04AD90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Slovakia

Available in:

Slovakia

Package description:

(ID2) 140 Film-coated tablet: unspecified outer container with 10 Blister (PolyVinyl Chloride; PolyVinylidene Chloride; Aluminium) each with 14 Film-coated tablet
(ID1) 56 Film-coated tablet: unspecified outer container with 4 Blister (PolyVinyl Chloride; PolyVinylidene Chloride; Aluminium) each with 14 Film-coated tablet

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:

Wirtschaftsgenossenschaft deutscher Tieraerzte eG

Marketing authorisation date:

28/06/2023

Manufacturing sites for batch release:

Artesan Pharma GmbH & Co. KG

Responsible authority:

Authorisation number:

96/007/MR/23-S

Date of authorisation status change:

28/06/2023

Reference member state:

Germany

Procedure number:

DE/V/0198/001

Concerned member states:

Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland
France Greece Hungary Ireland Italy Latvia Lithuania Luxembourg
Netherlands Norway Poland Portugal Romania Slovakia Slovenia Spain
Sweden United Kingdom (Northern Ireland)

To consult adverse reactions on veterinary medicinal products please go to
www.adrreports.eu/vet

Documents

Combined File of all Documents

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